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(71) Applicant (for all designated States except US): **GIVEN IMAGING LTD.** [IL/IL]; 2 HaCarmel St., Industrial Park, 20692 Yotneam (IL).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **LEWKOWIZC, Shlomo** [IL/IL]; 47 Habonim Street, 36031 Tivon (IL). **GAT, Daniel** [IL/IL]; 5 HaAarava Street, Nesher, 36863 Haifa (IL). **GLUKHOVSKY, Arkady** [IL/IL]; 24/5 Hanuriot Street, 36790 Nesher (IL). **KHAIT, Semion** [IL/IL]; 290/8 Mordot Tiberias St., 14350 Tiberias (IL).

(74) Agents: **EITAN, PEARL, LATZER & CO-HEN-ZEDEK** et al.; 2 Gav Yam Center, 7 Shenkar Street, 46725 Herzlia (IL).

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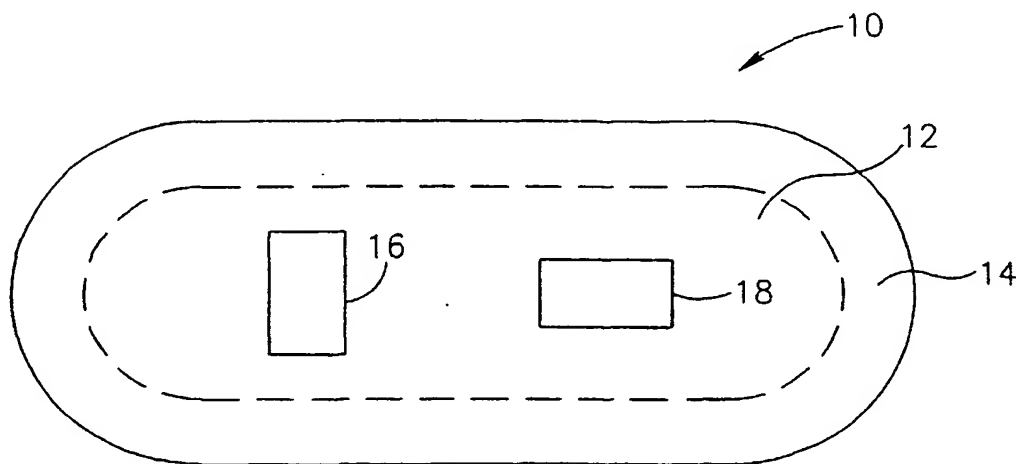
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(54) Title: **A FLOATABLE IN VIVO SENSING DEVICE**



(57) Abstract: The invention relates to an in vivo sensing device having a specific gravity of about 1 or a volume to weight ratio that enables it essentially to float. In one embodiment the in vivo sensing device consists of an image sensor system and a buoyant body. The buoyant body, which is attached to the sensor system or which can optionally house one or more elements of the sensor system, keeps the sensor system essentially floating in a body lumen liquid.

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## A FLOATABLE IN VIVO SENSING DEVICE

### FIELD OF THE INVENTION

The present invention relates to the field of in vivo sensing. Specifically, the  
5 present invention relates to a floatable in vivo sensing device that can be carried by liquid.

### BACKGROUND OF THE INVENTION

In vivo sensors are non invasive tools used in diagnosis of diverse body  
10 systems. For example, ingestible devices may be used for sensing in vivo conditions in the gastrointestinal (GI) tract, such as, temperature, pH or pressure. Ingestible imaging devices can be used for imaging the gastrointestinal tract. For example, a capsule comprising a sensor, such as an image sensor, may be ingested and moved passively through the small intestine by peristalsis while imaging or otherwise  
15 sensing the small intestine. However, passive movement of objects through larger body lumens, such as the large intestine, may be slower and unpredictable. The large intestine, or colon, whose main function is to remove much of the water from the stool and to store the stool, begins with the cecum, a small saclike evagination, then continues with the ascending colon, from the appendix in right groin up to a  
20 flexure at the liver, transverse colon, liver to spleen, descending colon, spleen to left

groin, then sigmoid (S-shaped) colon back to midline and anus. The colon has three longitudinal muscle bands whose actions assist movement through the colon.

It is sometimes advantageous to move objects through the colon independently of the natural action of the colon muscles. For example, delivery of a medicament to a specific location in the colon may be time dependant and cannot  
5 rely on the natural movement in the colon. Also a device for imaging the colon might benefit from being actively moved through the colon so as to efficiently view the colon.

Current methods of moving objects, especially imaging devices, through  
10 the colon involve the use of endoscopes, typically colonoscopes, which are expensive and inconvenient for patient use, and do not always enable to reach distal parts of the colon, such as the cecum and the right colon.

## 15 SUMMARY OF THE INVENTION

An embodiment of the present invention provides an essentially floatable sensing device that can be carried by liquid. The sensing device according to an embodiment of the invention is useful in sensing, such as by imaging, lumens containing or capable of containing a bulk of liquid. One example of such a lumen is  
20 the large intestine. The essentially floatable sensing device of the invention is carried by the liquid and can thus be moved through the lumen together with the bulk of liquid.

According to one embodiment of the invention the in vivo sensing device has a specific gravity (SG) of about 1.0 or a volume to weight ratio that enables it essentially to float.

In one embodiment the in vivo sensing device comprises a sensor system  
5 and a buoyant body. The buoyant body, which can optionally house one or more elements of the sensor system, may have a specific gravity of about 1.0, which is just about floating, or a volume to weight ratio that enables it essentially to float in a body lumen liquid. In another embodiment of the invention the buoyant body is a buoy attached to a sensor system, which keeps the sensor system essentially floating  
10 in a body lumen liquid.

The sensor system may comprise any sensor suitable for sensing in vivo environment parameters, such as pH, temperature, conductivity, shear, pressure and so on. The sensor system can include all the elements necessary for in vivo sensing, as known in the art. In one embodiment the sensor is an image sensor.

15 According to a further embodiment of the invention there is provided a method for sensing body lumens, such as the stomach or large intestine. Typically, the method involves moving an in vivo sensor through the body lumen. The method, according to one embodiment of the invention, comprises the steps of providing an essentially buoyant sensor system and sensing the body lumen.  
20 Preferably the body lumen contains a bulk of liquid. In an embodiment of the invention the method is used for sensing a patient's large intestines and may comprise the steps of clearing the large intestine of its contents, for example by

using standard laxatives and prep. solutions and filling the large intestine with a liquid.

According to yet another embodiment of the invention high osmotic pressure compositions, such as contrast agents, are utilized to assist in the  
5 movement of objects through the colon.

## BRIEF DESCRIPTION OF THE DRAWINGS

The present invention will be understood and appreciated more fully from the following detailed description taken in conjunction with the drawings in which:

5           Figure 1 is a schematic illustration of a floatable sensing device in accordance with an embodiment of the invention;

          Figure 2A is a schematic illustration of a floatable sensing device in accordance with another embodiment of the invention;

          Figure 2B is a schematic illustration of a floatable sensing device with a  
10   packaged buoy in accordance with an embodiment of the invention; and

          Figures 3A and 3B are schematic illustrations of a floatable sensing device in accordance with yet other embodiments of the invention.

15

## DETAILED DESCRIPTION OF THE INVENTION

A sensing device, according to an embodiment of the invention, is schematically illustrated in Fig 1. The sensing device 10 is an autonomous capsule shaped device and can thus be easily moved through the small intestine. However, it should be appreciated that an autonomous sensing device according to different  
5 embodiments of the invention can have any shape or design suitable for being inserted and moved through body lumens, such as the gastrointestinal tract, the reproductive system, blood vessels, etc.

Device 10 includes a sensor system 12 and a buoyant body 14. In one  
10 embodiment, the buoyant body 14 is essentially a part of the sensor system 12. The sensor system 12 may be a pH sensing system as known in the art, an image sensor as known in the art or other known in vivo sensing systems such as an in vivo temperature measuring system, a pressure sensor, a shear sensor, a sensor of electrical conductivity and other known in vivo sensors. In other embodiments a  
15 combination of sensors may be used. In one embodiment of the invention the sensor system 12 may be an imaging device comprising, for example, at least one image sensor (such as image sensor 16), one or more illumination source (not shown) and one or more transmitter for transmitting image signals to an external receiving system (not shown). In vivo imaging systems that can be utilized with the present  
20 invention are described, for example, with reference to Fig. 2A below, or in WO 01/65995 or in US Patent Number 5,604,531. In another embodiment the sensor system 12 includes a temperature measuring system. A temperature measuring

system according to an embodiment of the invention may comprise an image sensor 16 having an image sensing module in communication with an integrating unit (not shown) for detecting the dark current of the image sensor image sensing module and for calculating the temperature of the image sensor. The integrating unit further  
5 calculates the temperature of the environment or the temperature of the environment may be calculated, based on data from the integrating unit, by a separate unit that is in communication with the integrating unit. A temperature measuring system that may be utilized according to an embodiment of the invention is described, for example, in WO 01/10291.

10 Typically, the sensor system 12 is powered by a battery 18, however other embodiments are possible, for example, the sensor system may be power induced by an external electromagnetic field.

The buoyant body 14 lends a volume to weight ratio, or a specific gravity to the device 10 that enables it to float in a liquid filled lumen.

15 SG - specific gravity - is  $\text{weight [gr.] / weight of water in the amount displaced by the device 10 (when reaching maximal volume) or for sealed objects, SG is weight [gr] / volume [cubic cm]}$ . Typically, buoyant body 14 can be filled with a substance lighter than the body lumen liquid, such as gaseous  $\text{CO}_2$ ,  $\text{O}_2$  or air.

In another embodiment of the invention schematically illustrated in Fig.2A,  
20 the sensor system is an image sensor system 22 and the buoyant body is an inflatable buoy 24. The buoy 24 may be packaged such that it is not buoyant while in packaging. However, release of the buoy from its packaging lends buoyancy to



the sensor system. The buoy may be released from its packaging at a desired location or point in time, such that the sensor system 22 may acquire buoyancy according to specific requirements. For example, the floatable sensing device according to an embodiment of the invention may be ingested and moved by  
5 peristalsis through the small intestine while its buoy is packaged. When the device enters the large intestine the buoy is released from its packaging and the device can then float in the bulk of liquid in the large intestine and be carried by the bulk of liquid to all areas of the large intestine. Thus, the device is moved through the large intestine, effectively sensing the lumen. The mechanisms by which the buoy is  
10 released from its packaging can be externally controlled or automatically controlled as will be further described below.

The image sensor system 22 includes an image sensor 26, for example a charge coupled device (CCD) or a complementary metal oxide semiconductor (CMOS) image sensor, illumination sources 23, an optical system which includes an  
15 optical window 21 through which the body lumen is illuminated and imaged and lenses and/or mirrors (not shown) for collimating light, a transmitter 25 for transmitting image signals to an external receiving system (not shown) and a battery 27 for supplying power to the elements of the image sensor system 22.

The inflatable buoy 24 is a typically elastic compartment containing air,  
20 which is attached to the image sensor system 22. The inflatable buoy 24 will lend buoyancy to the image sensor system 22 such that it can essentially float in the

liquid filling a body lumen and be thus moved through the body lumen, obtaining images of essentially all parts of the body lumen.

In another embodiment of the invention, schematically illustrated in Fig. 2B an image sensor system 22, as above, comprises an inflatable buoy 24'. The inflatable buoy 24' is rolled and packaged to a small size. In its packaged form inflatable buoy 24' typically does not lend buoyancy to the image sensor system 22. The inflatable buoy 24' may be released from its packaging by a manually or automatically controlled mechanism, at a desired location *in vivo*. For example, the inflatable buoy 24' may contain gas releasing granules such as crystalline sodium bicarbonate, E-Z GasII effervescent granules by EZEM of NY, USA or similar oxygen releasing granules. Typically, these granules release gas (such as CO<sub>2</sub> or oxygen) upon contacting liquid. In an embodiment of the invention inflatable buoy 24' contains two compartments (not shown), one compartment containing gas releasing granules (for example 100 mg of granules) and the other containing a drop of liquid (for example 0.1 cc of water or saline). The compartments are kept separate while the inflatable buoy 24' is in its packaged form. Once the packaging is opened, the two compartments are merged and the drop of liquid contacts the gas releasing granules. Gas is released into the now unpacked buoy inflating the buoy and enhancing buoyancy.

The packaging can be achieved by a manually controlled mechanism, such as a bimorph material mechanism that could change configuration in accordance with controllable conditions, such as a temperature or electric voltage gradient, as

known in the art. Alternatively, the packaging mechanism can be time dependent or dependant on in vivo conditions such as pH or enzymatic activity. For example, packaging can be effected by a hydrocarbon such as a gelatin capsule which encases the image sensor system 22 and the inflatable buoy 24' and keeps the inflatable buoy 24' in a packaged form. According to one embodiment, the gelatin capsule slowly dissolves in the liquid present in the stomach, thereby releasing the inflatable buoy. The gelatin capsule, such as gelatin capsules provided by Capsugel USA, can be made to dissolve at a specific location along the GI tract, as known in the field of sustained release mechanisms. Thus, a device comprising image sensor system 22 and inflatable buoy 24' can be encased in a hydrocarbon capsule while moving through certain parts of a patient's GI tract and free of the encapsulation in other parts of the GI tract. While free of the encapsulation, inflatable buoy 24' will be inflated and the device will be able to essentially float in the GI tract liquid. In another embodiment the packaging can be effected by degradable sutures known in the art, such that the packaging of the inflatable buoy 24' is released when the suture is degraded.

Another embodiment of the invention is schematically illustrated in Figs. 3A and 3B, in which a sensor system 32, such as an imaging sensor system or other sensing systems, for example, those described above, is attached to a buoy 34, which is typically a light vessel made of, for example, plastic such as isoplast, filled with a substance lighter than the body lumen liquid. In certain embodiments of the

invention the buoy 34 can contain accessory components or materials, such as additional batteries 31.

The sensor system 32 and the buoy 34 can be attached by a flexible sleeve 33 (Fig. 3A), which, if filled with a substance lighter than the body lumen liquid, can also assist in enhancing the buoyancy of the sensor system 32. Alternatively, the sensor system 32 and the buoy 34 can be attached by a wire 35 (Fig. 3B). The wire 35 can couple sensor system 32 and the buoy 34, for example, if electrical coupling is needed for utilizing the additional batteries 31 by the sensor system 32.

According to a further embodiment of the present invention there is provided a method for sensing body lumens, such as the stomach or large intestine. In an embodiment of the invention a floatable sensing device, such as the devices described above, is inserted in vivo. The sensing device may include a sensor system and a buoyant body. The buoyant body may be inflated prior to being inserted in vivo or may be inflated at a specific location or point in time while in a patient's body, as described above. Preferably, the sensor system and buoyant body are placed in a body lumen containing a bulk of liquid. More preferably there is a flow of liquids through the body lumen, upon which the sensor system is carried through the body lumen. Thus, this method can be used, in one embodiment, for imaging or otherwise sensing a patient's large intestines. In one embodiment the patient's large intestine is initially cleared of its contents, for example by inducing bowel movement by administering a laxative or an enema. Further, the patient's large intestine is filled with a liquid, for example by drinking high osmolarity

liquids, which retain liquids within the large intestine for longer periods. Typically, a liquid loaded intestine and/or additional laxatives, administered while the sensor system is in the large intestine, will induce bowel movement and cause a flow of the bulk of liquids in the large intestine. The induced flow will enhance movement of the sensor system through the large intestine, thereby facilitating sensing of most areas of the lumen.

According to one embodiment of the invention an imaging device for imaging the GI tract can be moved through the large intestine by utilizing, for example, a high osmotic pressure composition that is essentially not absorbed by the intestine. According to other embodiments an object other than an imaging device, for example a device for sustained release of medicaments to the colon, can be moved through the large intestine by utilizing a high osmotic pressure composition that is essentially not absorbed by the intestine. Objects moved through the large intestine utilizing a high osmotic pressure composition according to an embodiment of the invention may or may not be floatable.

A wireless in vivo imaging device such as the sensor systems described above or a device including a compartment for collecting or distributing substances from or to the GI tract, such as the device described in WO00/22975, can be moved through the large intestine according to an embodiment of the invention such that it can be used for diagnostic and/or for therapeutic purposes.

In one embodiment, an in vivo imaging device is introduced into the GI tract by utilizing a colonoscope. In another embodiment the in vivo imaging device

is ingested by a patient and traverses the small intestine pushed along by natural peristalsis. When the device reaches the cecum it may remain in the cecum for typically long periods of time. According to an embodiment of the invention a high osmotic pressure composition that is essentially not absorbed by the intestine, such as a contrast agent, may be administered to the patient. The high osmotic pressure composition typically progresses through the small intestine arriving at the cecum. The composition usually progresses in the intestine faster than the device, pushing in its progression the device through the different parts of the colon. An example of a high osmotic pressure composition that may be used according to an embodiment of the invention is barium sulfate or gastrografin. The osmotic pressure of gastrografin at 37°C is 55.1 Atm and its osmolality is 2.15 (osm/kg H<sub>2</sub>O).

A wireless ingestible imaging capsule as described above was administered under Helsinki Committee guidelines to normal healthy volunteers in the standard fashion. The volunteers underwent standard colonoscopic preparation consisting of a 24-hour liquid diet and 4 ounces of Fleet phosphasoda the evening before and the morning of the procedure. The volunteers were encouraged to drink a large volume of fluid before and after ingestion of the capsule. The passage of the capsule was monitored on-line. The volunteers were given 8 ounces of gastrografin diluted as for regular body CT examination [5%], every 15 minutes starting 2.5 hours after ingestion of the capsule, up to a total of 2 liters. The volunteers were allowed to eat a normal meal 4 hours after ingestion of the capsule.

**Results:** In one case, the gastric emptying time was 42 minutes, and the capsule reached the cecum after 4 hours and 16 minutes. The capsule was excreted from the colon 22 hours after ingestion. Images of various parts of the colon were acquired for more than 3 hours. In the second case, gastric emptying occurred after 5 8 minutes, and the capsule reached the cecum after 8 hours. Images of all parts of the colon were acquired for 2.5 hours. The capsule was excreted after 10.5 hours, still working.

**Conclusion:** The first successful passage through the right colon in the first case and the entire colon in the second case of a functioning imaging capsule using a 10 novel technique is described. The use of gastrografin was based on clinical observation in GI radiological procedures, such as CT. This agent may have an important role in the movement of the imaging capsule through the colon.

Gastrografin contains a mixture of sodium amidotrizoate and meglumine amidotrizoate in a proportion of 10:66 (amidotrizoic acid or diatrizoic acid: 15 3,5-bis-acetamido-2,4,6-triiodobenzoic acid). 1 mL gastrografin contains sodium amidotrizoate 100,00 mg and meglumine amidotrizoate 660,00 mg (sodium diatrizoate and meglumine diatrizoate) in aqueous solution plus flavorings and a wetting agent. The contrast-giving substances in gastrografin are salts of amidotrizoic acid in which the X-ray absorbing iodine is present in stable chemical 20 bond. Following oral administration only about 3% of the amidotrizoic acid is absorbed by the stomach and intestines. This portion is eliminated mainly via the kidneys.

Gastrografin can be used either orally or as an enema. Gastrografin typically travels through the colon faster than the capsule and is a clear fluid, allowing a clear view of the colon. Another advantage of gastrografin is its opacity to x-ray, allowing direct view in fluoroscopy and enabling monitoring of the  
5 passage of the capsule through the GI tract.

It will be appreciated by persons skilled in the art that the present invention is not limited to what has been particularly shown and described hereinabove. Rather the scope of the present invention is defined only by the claims which follow:



## CLAIMS

1. An in vivo sensing device having a specific gravity of about 1.
2. An in vivo sensing device having a volume to weight ratio that enables said sensing device essentially to float in a body lumen liquid.
- 5 3. The in vivo sensing device according to claim 1 or 2 comprising at least one in vivo sensor.
4. The in vivo sensing device according to claim 3 wherein the in vivo sensor is an image sensor.
5. The in vivo sensing device according to claim 3 wherein the in vivo sensor is  
10 selected from the group consisting of: an image sensor, a pH meter, a temperature sensor, a pressure sensor, a shear sensor, a sensor of electrical conductivity or a combination thereof.
6. The in vivo sensing device according to claim 4 further comprising at least one illumination source.
- 15 7. The in vivo sensing device according to claim 4 further comprising at least one transmitter for transmitting image data to an external receiving system.
8. The in vivo sensing device according to claim 1 or 2 comprising a sensor system and a buoyant body.
9. The in vivo sensing device according to claim 8 wherein the buoyant body  
20 houses the sensor system.

10. The in vivo sensing device according to claim 8 wherein the buoyant body is attached to the sensor system.
11. The in vivo sensing device according to claim 8 comprising at least one in vivo sensor.
- 5 12. The in vivo sensing device according to claim 11 wherein the in vivo sensor is an image sensor.
13. The in vivo sensing device according to claim 11 wherein the in vivo sensor is selected from the group consisting of: an image sensor, a pH meter, a temperature sensor, a pressure sensor, a shear sensor, a sensor of electrical  
10 conductivity or a combination thereof.
14. The in vivo sensing device according to claim 12 further comprising at least one illumination source.
15. The in vivo sensing device according to claim 12 further comprising at least one transmitter for transmitting image data to an external receiving system.
- 15 16. The in vivo sensing device according to claim 9 comprising at least one image sensor and at least one transmitter.
17. The in vivo sensing device according to claim 8 wherein the buoyant body is inflatable.
18. The in vivo sensing device according to claim 17 wherein the buoyant body  
20 comprises gas releasing granules.

19. The in vivo sensing device according to claim 17 wherein the buoyant body comprises a plurality of compartments, at least one compartment comprising gas releasing granules and at least another compartment containing liquid.

20. A method for sensing a body lumen comprising the step of moving an in vivo  
5 sensing device according to claim 1 or 2, through a body lumen.

21. A method for sensing a large intestine comprising the steps of:

clearing the large intestine of contents;

filling the large intestine with a liquid; and

10 providing the large intestine with an in vivo sensing device  
according to claim 1 or 2 or 8.

22. A method for moving an object through a large intestine comprising the steps  
of:

placing an object in a gastrointestinal tract; and

15 administering to the gastrointestinal tract a suitable amount of a  
high osmotic pressure composition that is essentially not absorbed by the  
intestine.

23. The method according to claim 22 further comprising a step of performing a  
colonoscopic preparation prior to the step of placing the object.

24. The method according to claim 22 wherein placing an object in a  
20 gastrointestinal tract comprises placing the object in the mouth, esophagus or  
stomach.

25.The method according to claim 22 wherein the step of placing an object is performed by ingesting the object.

26.The method according to claim 22 wherein the step of placing an object is performed by utilizing a colonoscope.

5 27.The method according to claim 22 wherein the object is a diagnostic or therapeutic device or a device that is both diagnostic and therapeutic for monitoring or treating or for monitoring and treating the gastrointestinal tract.

28.The method according to claim 22 wherein the object is a device for sustained release of medicaments to the colon.

10 29.The method according to claim 22 wherein the object is an in vivo imaging device.

30.The method according to claim 22 wherein the high osmotic pressure composition comprises a contrasting agent.

31.The method according to claim 22 wherein the high osmotic pressure  
15 composition comprises gastrografin or barium sulfate.

32.The method according to claim 22 wherein the step of administering a high osmotic pressure composition comprises administering 8 ounces of gastrografin diluted to about 5%, in predetermined time intervals after placing of the object, up to a total of 2 liters.

20 33.The method according to claim 32 wherein the predetrmined time intervals are every 15 minutes starting from about 2.5 hours after placing of the object.

34. Use of contrasting agents in the preparation of a high osmotic pressure composition that is essentially not absorbed by the intestine for moving an object in a colon.
35. The use according to claim 34 wherein the contrasting agent is gastrografin or  
5       barium sulfate.
36. The use according to claim 34 wherein the object is a diagnostic or therapeutic device or a device that is both diagnostic and therapeutic for monitoring or treating or for monitoring and treating the gastrointestinal tract.
37. The use according to claim 34 wherein the object is a device for sustained  
10       release of medicaments to the colon.
38. The use according to claim 34 wherein the object is an in vivo imaging device.
39. A high osmotic pressure composition that is essentially not absorbed by the intestine for moving a sensor system through a patient's colon comprising a  
15       contrasting agent and additives.
40. A composition according to claim 39 wherein the contrasting agent is gastrografin or barium sulfate.
41. A composition according to claim 39 wherein the additives comprise flavorings and/or wetting agents.
- 20    42. A method for wireless imaging of the colon comprising the steps of  
          placing a wireless in vivo imaging device in a gastrointestinal tract;

administering to the gastrointestinal tract a suitable amount of a high osmotic pressure composition that is essentially not absorbed by the intestine; and

activating the in vivo imaging device to obtain images of the colon.

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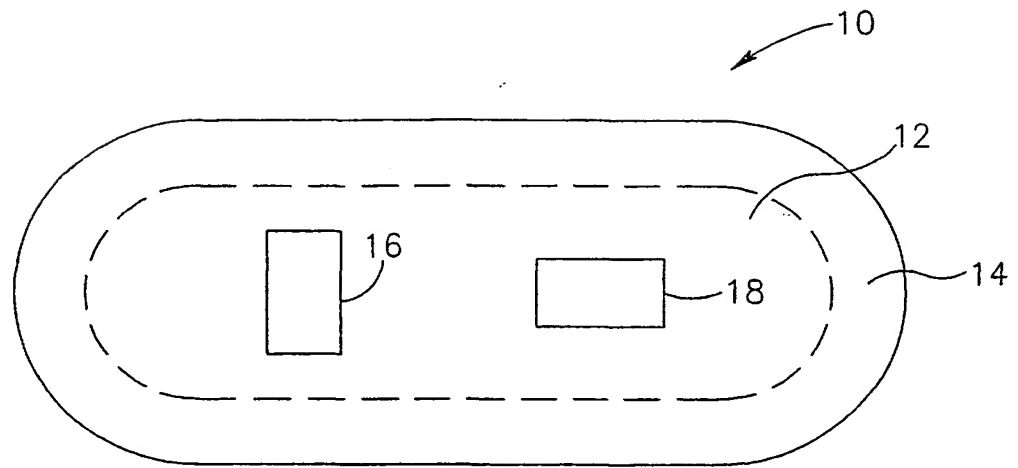


FIG. 1

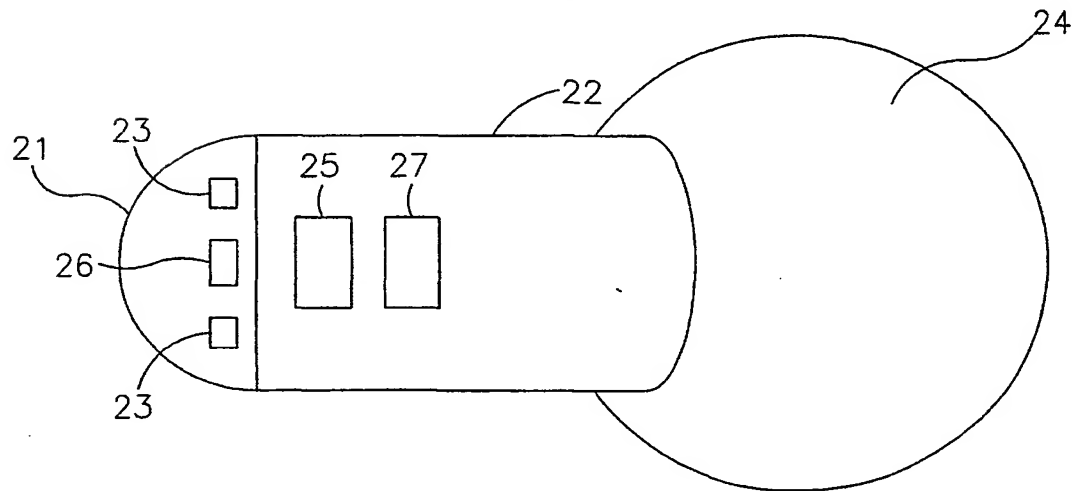


FIG. 2A

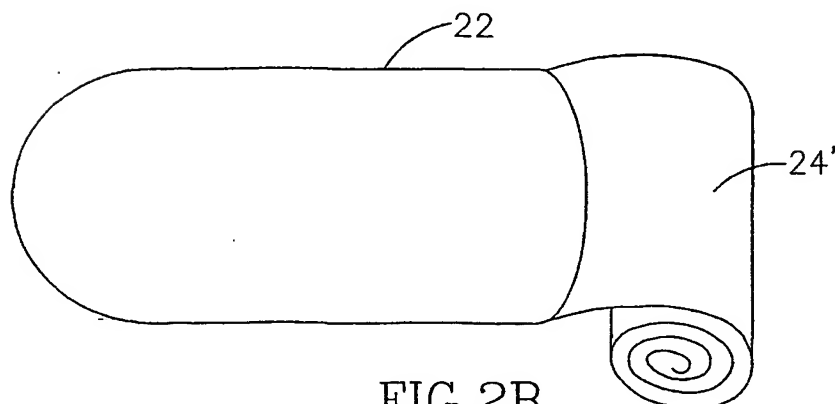


FIG. 2B

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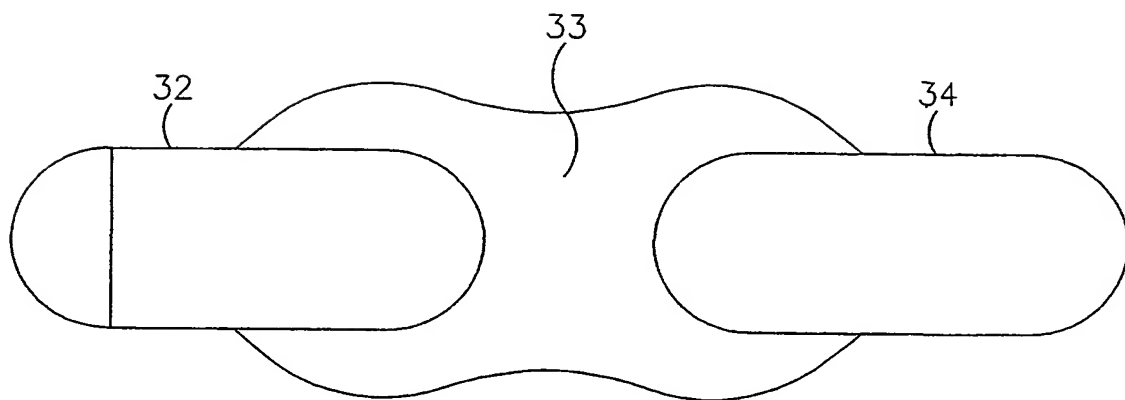


FIG. 3A

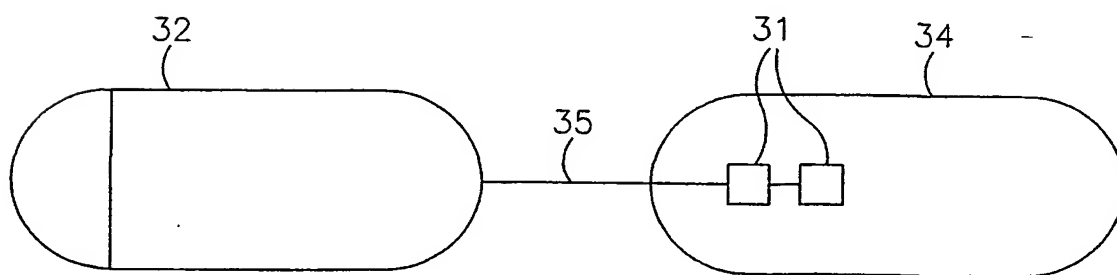


FIG. 3B

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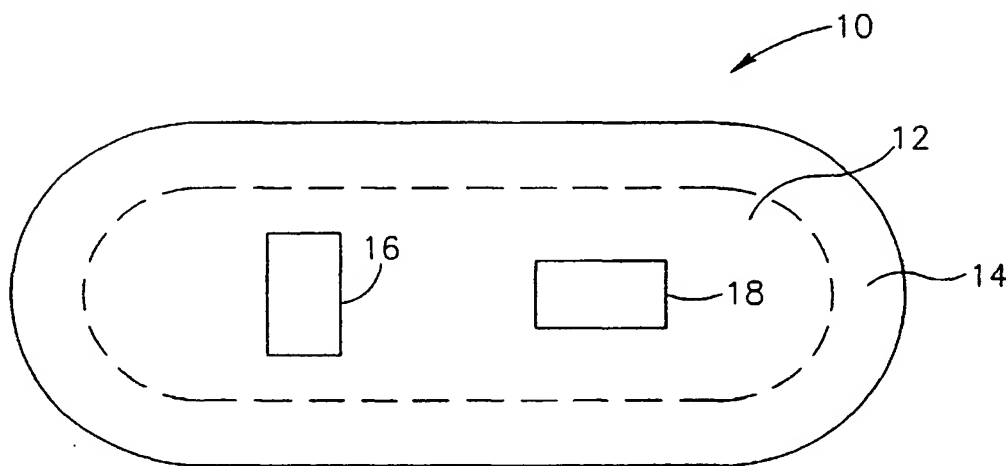
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- (71) Applicant (*for all designated States except US*): **GIVEN IMAGING LTD.** [IL/IL]; 2 HaCarmel St., Industrial Park, 20692 Yoqneam (IL).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): **LEWKOWIZC, Shlomo** [IL/IL]; 47 Habonim Street, 36031 Tivon (IL). **GAT, Daniel** [IL/IL]; 5 HaAarava Street, Neshet, 36863 Haifa (IL). **GLUKHOVSKY, Arkady** [IL/IL]; 24/5 Hanuriot Street, 36790 Neshet (IL). **KHAIT, Semion** [IL/IL]; 290/8 Mordot Tiberias St., 14350 Tiberias (IL).
- (74) Agents: **EITAN, PEARL, LATZER & CO-HEN-ZEDEK** et al.; 2 Gav Yam Center, 7 Shenkar Street, 46725 Herzlia (IL).
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- Published:  
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[Continued on next page]

(54) Title: A FLOATABLE IN VIVO SENSING DEVICE



(57) Abstract: An in vivo sensing device has a specific gravity of about 1 or a volume to weight ratio that enables it essentially to float. In one embodiment the in vivo sensing device (10) consists of an image sensor system (12) and a buoyant body (14). The buoyant body, which is attached to the sensor system or which can optionally house one or more elements (16, 18) of the sensor system, keeps the sensor system essentially floating in a body lumen liquid.

WO 02/095351 A3



*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/IL02/00392

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61B 5/00, 5/103, 5/117

US CL : 600/593

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 600/101, 109, 301, 410, 411, 412, 431, 481, 505, 547, 549, 561, 587, 593; 348/65, 77; 128/898, 903; 455/66, 95, 100

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
Please See Continuation Sheet

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category *        | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No.                 |
|-------------------|--|---------------------------------------|
| X,P<br>---<br>A,P | US 6,240,312 B1 (ALFANO et al) 29 May 2001, see entire document.                   | 1-16,20<br>-----<br>17-19,21          |
| X<br>---<br>A     | US 5,604,531 A (IDDAN et al) 18 February 1997, see entire document.                | 1-16,20,21<br>-----<br>17-19,22-38,42 |
| X<br>---<br>A     | US 6,149,581 A (KLINGENSTEIN) 21 November 2000, see entire document.               | 22-31,34-40,42<br>-----<br>32-33,41   |
| A                 | US 6,099,482 A (BRUNE et al) 08 August 2000, see entire document.                  | 1-22,42                               |

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

|   |  |
|---|--|
| * Special categories of cited documents:  |  |
| "A" document defining the general state of the art which is not considered to be of particular relevance  | "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention  |
| "E" earlier application or patent published on or after the international filing date   | "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone   |
| "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) | "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art |
| "O" document referring to an oral disclosure, use, exhibition or other means  | "&" document member of the same patent family  |
| "P" document published prior to the international filing date but later than the priority date claimed  |  |

Date of the actual completion of the international search

24 October 2002 (24.10.2002)

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Name and mailing address of the ISA/US

Commissioner of Patents and Trademarks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703)305-3230

Authorized officer

Charles A. Marmor, II

Telephone No. (703) 308-0858

Form PCT/ISA/210 (second sheet) (July 1998)

# INTERNATIONAL SEARCH REPORT

PCT/IL02/00392

## Continuation of B. FIELDS SEARCHED Item 3:

EAST

search terms: pill, capsule, float, specific gravity, volume, weight, buoyant, sensor, transducer, pH, imaging, camera, pressure, temperature, shear, conductivity, intestinal, gastrointestinal, colon, contrasting agent, additive, wetting, flavoring, barium sulfate, gastrografin